

GenCore version 5.1.4.p5.4578  
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OM protein - protein search, using sw model

Run on: March 14, 2003, 09:22:34 ; Search time 24.9038 Seconds

(without alignments)  
2108.144 Million cell updates/sec

Title: US-09-836-077-4

Perfect score: 2120  
Sequence: 1 MTPPPGSAAPSAPRARVLS.....TFQVADSHPEVAQRPEWGP 394

Scoring table: BLOSUM62

Gapop 10.0, Gapext 0.5

Searched: 908470 seqs, 133250620 residues

Total number of hits satisfying chosen parameters: 908470

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database :

A.GeneSeq\_101002.\*  
1: /SIDSI/gcgdata/geneSeq/geneSeq-emb1/AA1980.DAT.\*  
2: /SIDSI/gcgdata/geneSeq/geneSeq-emb1/AA1981.DAT.\*  
3: /SIDSI/gcgdata/geneSeq/geneSeq-emb1/AA1982.DAT.\*  
4: /SIDSI/gcgdata/geneSeq/geneSeq-emb1/AA1983.DAT.\*  
5: /SIDSI/gcgdata/geneSeq/geneSeq-emb1/AA1984.DAT.\*  
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8: /SIDSI/gcgdata/geneSeq/geneSeq-emb1/AA1987.DAT.\*  
9: /SIDSI/gcgdata/geneSeq/geneSeq-emb1/AA1988.DAT.\*  
10: /SIDSI/gcgdata/geneSeq/geneSeq-emb1/AA1989.DAT.\*  
11: /SIDSI/gcgdata/geneSeq/geneSeq-emb1/AA1990.DAT.\*  
12: /SIDSI/gcgdata/geneSeq/geneSeq-emb1/AA1991.DAT.\*  
13: /SIDSI/gcgdata/geneSeq/geneSeq-emb1/AA1992.DAT.\*  
14: /SIDSI/gcgdata/geneSeq/geneSeq-emb1/AA1993.DAT.\*  
15: /SIDSI/gcgdata/geneSeq/geneSeq-emb1/AA1994.DAT.\*  
16: /SIDSI/gcgdata/geneSeq/geneSeq-emb1/AA1995.DAT.\*  
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19: /SIDSI/gcgdata/geneSeq/geneSeq-emb1/AA1998.DAT.\*  
20: /SIDSI/gcgdata/geneSeq/geneSeq-emb1/AA1999.DAT.\*  
21: /SIDSI/gcgdata/geneSeq/geneSeq-emb1/AA2000.DAT.\*  
22: /SIDSI/gcgdata/geneSeq/geneSeq-emb1/AA2001.DAT.\*  
23: /SIDSI/gcgdata/geneSeq/geneSeq-emb1/AA2002.DAT.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	2120	100.0	394	20	AAW92805
2	2115	99.8	664	21	AAV82434
3	1897	89.5	666	20	AAV39445
4	1897	89.5	666	20	AAV28552
5	1897	89.5	666	20	AAW92804
6	1897	89.5	666	21	AAV82433
7	1897	89.5	666	21	AAV55009
8	1897	89.5	666	22	AAE02650
9	1755	82.8	634	21	AAV56854
10	948.5	44.7	215	20	AAV28553

11	902	42.5	606	21	AAV56855
12	552	26.0	379	22	AAW93440
13	397	18.7	893	23	AAW97964
14	389.5	18.4	861	18	AAW17658
15	389.5	18.4	861	19	AAW58540
16	389.5	18.4	861	22	AAW81035
17	389.5	18.4	861	22	AAW51251
18	385.5	18.2	832	22	AAE03818
19	385.5	18.2	832	23	AAW64522
20	385.5	18.2	837	21	AAV99410
21	385.5	18.2	837	22	AAU29250
22	385.5	18.2	837	22	AAW66159
23	376.5	17.8	791	23	AAU77413
24	368	17.4	796	22	AAW62727
25	366	17.3	746	19	AAV21264
26	363	17.1	771	16	AAW71380
27	363	17.1	771	22	AAW62726
28	361.5	17.1	862	18	AAW17657
29	361.5	17.1	862	22	AAW81036
30	361.5	17.1	862	22	AAW51252
31	346	16.3	660	20	AAV13463
32	346	16.3	660	21	AAW28523
33	346	16.3	660	22	AAW70132
34	346	16.3	660	22	AAW31694
35	342.5	16.2	775	19	AAW63748
36	341.5	16.1	441	16	AAW71381
37	341	16.1	886	23	AAW97963
38	339.5	16.0	590	22	AAW48373
39	339.5	16.0	596	22	AAW48374
40	339.5	16.0	624	22	AAW48378
41	337.5	15.9	833	22	AAE03640
42	334.5	15.8	833	23	AAE18214
43	334.5	15.8	833	23	AAE18215
44	333.5	15.7	833	23	AAE18213
45	332.5	15.7	775	20	AAV43090

#### ALIGNMENTS

RESULT 1	AAW92805	standard; Protein; 394 AA.
AC	AAW92805;	
XX		
DT	07-MAY-1999 (first entry)	
XX		
DE	EP-892047 Seq ID 4.	
XX		
KW	Semaphorin L; human; immunosuppressant; anti-inflammatory; gene therapy;	
KW	organ transplantation; inflammation therapy; immunotherapy; agonist;	
KW	immunomodulatory; antagonist.	
XX		
OS	Homo sapiens.	
XX		
PN	EP892047-A2.	
XX		
PD	20-JAN-1999.	
XX		
PF	06-JUL-1998: 98EP-0112470.	
XX		
PR	11-FEB-1998: 98DE-1005371.	
PR	09-JUL-1997: 97DE-1029211.	
XX		
PI	(HMRI) HOECHST MARION ROUSSEL DEVU GMBH.	
XX		
XX	Ensser A, Fleckenstein B;	
DR	WPI; 1999-083564/08.	
XX		
PT	New semaphorin L proteins - used as immunosuppressants and anti-inflammatory agents in organ transplants, inflammation therapy,	

PT immunotherapy and gene therapy  
XX  
PS Claim 4; Page 64-65; 135pp; German.  
XX

CC This invention describes a novel human semaphorin 1 protein. This protein  
CC or its encoding DNA are useful as immunosuppressants and/or  
CC anti-inflammatory agents in organ transplantation, inflammation therapy,  
CC immunotherapy and gene therapy. The DNA can be used to produce knock-out  
CC or knock-in animals for research purposes. The proteins or DNA can be  
CC used to search for the corresponding receptors or to screen for  
CC immunomodulatory agonists or antagonists.  
XX

XX Sequence 394 AA:

Query Match 100.0%; Score 2120; DB 20; Length 394;  
Best Local Similarity 100.0%; Pred. No. 5,76-204;  
Matches 394; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MTPPPGPAAPAPARAVLSLPARFGLPLRLLLVFWVAASAGSHSGPRISAVMKG 60  
DB 1 MTPPPGPAAPAPARAVLSLPARFGLPLRLLLVFWVAASAGSHSGPRISAVMKG 60  
QY 61 QDHVDFSOPEPHTVLFHEPGSFSVWVGGRKAYHFNFPREGKASVTVNIGSTKSGCDK 120  
DB 61 QDHVDFSOPEPHTVLFHEPGSFSVWVGGRKAYHFNFPREGKASVTVNIGSTKSGCDK 120  
QY 121 QDCGNVITLLERRGNGLVCGTNARKPCSCWNLVNDVSVSLGEMKGYAFSPDENSLVLF 180  
DB 121 QDCGNVITLLERRGNGLVCGTNARKPCSCWNLVNDVSVSLGEMKGYAFSPDENSLVLF 180  
QY 181 EGDVEYSTRKOEYNGKIPFRFRIGESLTYSDTVMNPQFIKATIVHQDAYDKITY 240  
DB 181 EGDVEYSTRKOEYNGKIPFRFRIGESLTYSDTVMNPQFIKATIVHQDAYDKITY 240  
QY 241 FFRENPNKPNPAPLVNSVAQLCRDGGESSLSYSKNTFLKAMLVCSDAATNENFR 300  
DB 241 FFRENPNKPNPAPLVNSVAQLCRDGGESSLSYSKNTFLKAMLVCSDAATNENFR 300  
QY 301 LODVFLPDPSCQMDRTRYGVFSNPWNYSAYCVYSLGIDIDRVFTSSLKGYHMGLSNPR 360  
DB 301 LODVFLPDPSCQMDRTRYGVFSNPWNYSAYCVYSLGIDIDRVFTSSLKGYHMGLSNPR 360  
QY 361 PGMCLPKKQPIPTETFOVADSHPEVAQRYEPMGP 394  
DB 361 PGMCLPKKQPIPTETFOVADSHPEVAQRYEPMGP 394

RESULT 2  
AAy82434  
ID AAY82434 standard; Protein; 664 AA.  
XX  
AC AAY82434;  
XX  
DT 27-JUN-2000 (first entry)  
XX  
DE Mouse CDw108 protein SEQ ID NO:8.  
XX  
KW Mouse; CDw108; detection; diagnosis; HIV; infection; anti-HIV.  
XX  
OS Mus musculus.  
XX  
FN WO200012700-A1.  
XX  
PD 09-MAR-2000.  
XX  
PE 25-AUG-1999; 99WO-JP04571.  
XX  
PR 26-AUG-1998; 98JP-0239687.  
XX  
PA (SHIO ) SHIONOGI & CO LTD.  
XX  
PI Yamada A, Kubo K, Itoh K;  
XX

DR WPI: 2000-246752/21.  
XX N-PSDB: AAA08189.  
XX

PT New CDw108 protein, useful in diagnosis of and as remedy for  
PT CDw108-associated diseases e.g. HIV-1 infection, and in study of  
PT biological functions and molecular specificity of CDw108 -  
XX  
XX Example 8; Page 64-69; 73pp; Japanese.  
XX

CC The present invention describes human CDw108. The CDw109 nucleotide  
CC and protein sequences can be used in the diagnosis and treatment of  
CC CDw108-associated diseases e.g. HIV-1 infection, and in study of  
CC biological functions and molecular specificity of CDw108. The present  
CC sequence represents mouse CDw108 given in an example from the present  
CC invention.  
XX

XX Sequence 664 AA:

Query Match 99.8%; Score 2115; DB 21; Length 664;  
Best Local Similarity 99.7%; Pred. No. 4,2e-203;  
Matches 393; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 MTPPPGPAAPAPARAVLSLPARFGLPLRLLLVFWVAASAGSHSGPRISAVMKG 60  
DB 1 MTPPPGPAAPAPARAVLSLPARFGLPLRLLLVFWVAASAGSHSGPRISAVMKG 60  
QY 61 QDHVDFSOPEPHTVLFHEPGSFSVWVGGRKAYHFNFPREGKASVTVNIGSTKSGCDK 120  
DB 61 QDHVDFSOPEPHTVLFHEPGSFSVWVGGRKAYHFNFPREGKASVTVNIGSTKSGCDK 120  
QY 121 QDCGNVITLLERRGNGLVCGTNARKPCSCWNLVNDVSVSLGEMKGYAFSPDENSLVLF 180  
DB 121 QDCGNVITLLERRGNGLVCGTNARKPCSCWNLVNDVSVSLGEMKGYAFSPDENSLVLF 180  
QY 181 EGDVEYSTRKOEYNGKIPFRFRIGESLTYSDTVMNPQFIKATIVHQDAYDKITY 240  
DB 181 EGDVEYSTRKOEYNGKIPFRFRIGESLTYSDTVMNPQFIKATIVHQDAYDKITY 240  
QY 241 FFRENPNKPNPAPLVNSVAQLCRDGGESSLSYSKNTFLKAMLVCSDAATNENFR 300  
DB 241 FFRENPNKPNPAPLVNSVAQLCRDGGESSLSYSKNTFLKAMLVCSDAATNENFR 300  
QY 301 LODVFLPDPSCQMDRTRYGVFSNPWNYSAYCVYSLGIDIDRVFTSSLKGYHMGLSNPR 360  
DB 301 LODVFLPDPSCQMDRTRYGVFSNPWNYSAYCVYSLGIDIDRVFTSSLKGYHMGLSNPR 360  
QY 361 PGMCLPKKQPIPTETFOVADSHPEVAQRYEPMGP 394  
DB 361 PGMCLPKKQPIPTETFOVADSHPEVAQRYEPMGP 394

RESULT 3  
AAy39445  
ID AAY39445 standard; Protein; 666 AA.  
XX  
AC AAY39445;  
XX  
DT 01-DEC-1999 (first entry)  
XX  
DE Human semaphorin 7.  
XX  
KW Semaphorin; transmembrane; secreted; neuroregeneration;  
XX immunosuppression; diabetes; multiple sclerosis; rheumatoid arthritis;  
XX proliferation; differentiation.  
XX  
OS Homo sapiens.  
XX  
FH Key  
FT Domain  
XX  
XX W09945114-A2.

PD		10-SEP-1999.	
XX	PX		
XX	PX	03-MAR-1999;	99WO-US04758.
XX	PR	03-MAR-1998;	98US-0076611.
XX	PA	(ZIMO ) ZYMOGENETICS INC.	
XX	PI	Holloway JL, Lofton-Day CE;	
DR	XX	WPI: 1999-540845/45.	
XX	DR	N-PSDB; AAZ20985, AAZ20986.	
PT	PT	New isolated human semaphorin ZSMF-7 polypeptides, used to develop products for treating e.g. immunodeficiencies, autoimmune diseases, inflammation, graft rejection and infective diseases -	
PS	PS	Claim 6: Page 101-102; 124pp; English.	
CC	CC	This sequence represents human ZSMF-7 semaphorin. The cDNA was isolated and amplified from a human testis cDNA library using PCR primers zc16189 (AAZ20989) and zc16188 (AAZ20990) which had been designed based upon an incomplete clone obtained from a human placenta library. Semaphorins have a variety of roles. They influence the direction and degree of axon and dendrite growth in nervous tissue, and may thus be useful as therapeutic agents for various neurodegenerative conditions. They are active in defining and directing development of various tissues and organs including those associated with muscle, fibroblasts, reproductive, endocrine and lymphatic tissues. ZSMF-7 plays a role as a mediator of immunosuppression, in particularly the activation and regulation of T lymphocytes. ZSMF-7 polypeptides would be useful additions to therapies for treating immunodeficiencies. ZSMF-7 is expressed in activated lymphocytes (MRL cells) and not in resting lymphocyte cells (CD4+ and CD8+) suggesting that it would be a useful tool for diagnosis and treatment of conditions where selective elimination of inappropriately activated T cells would be beneficial, such as in autoimmune diseases. In particular insulin dependent diabetes mellitus, Rheumatoid arthritis and multiple sclerosis. ZSMF-7 polypeptides can be used in vivo as anti-inflammatory agents,	
CC	CC	for inhibition of antigen in humoral and cellular immunity and for immunosuppression in graft and organ transplants.	
XX	SO	Sequence 666 AA;	
QY		Query Match 89.5%; Score 1897; DB 20: Length 666; Best Local Similarity 90.2%; Pred. No. 3.3e-181;	
Matches 358:	Conservative 8; Mismatches 27; Indels 4; Gaps 2		
OY	1	MTPPPPGGAAPSAPRARVLSLPAREGLPLRLLLFWMAAASAGHSNSGPRIASVWK-	59
Db	1	MTTPPPGGRAAPSAPEARVGVGPARRLGLPLRLRLTLMAAASAOGHLSGPRIFAWK	60
OY	60	--GDHVDSOEPEPHVLNHPGPSFVMWGGRKKVYNFMRECKNASVTYVNGSRKGC	117
Db	61	HVGODVDVDEGQTEPHVLEHEPGSSVWMWGGRKVLLPFRPECKNASVRYVNGSRKGC	120
OY	118	ODKOCGNATITLERRGNGLVCGTARKPCSGCNLVDSVSLGEMKGAPSPDENSL	177
Db	121	LDKRCOCENTILLERSSEBLACGTNAHRPSCMNLVNGIVV-PLGEMRGCAFPSPDENSL	179
OY	178	VLEPEDEVYSTIRKOENYNGKIPRFRRIRGESELVTSDTVMQNQFIKATIVHODAAYDDK	237
Db	180	VLEFEDEVYSTIRKOENYNGKIPRFRRIRGESELYTSDTVMQNPQFIKATIVHODAAYDDK	239
OY	238	IYFFREDNDPKNNPEAPLNVSVRAOQLCRDODGEGSSLVSKNWTFILAKMLYCDAATNRN	297
Db	240	IYFFREDNDPKNNPEAPLNVSVRAOQLCRDODGEGESLSVKWNWTFILAKMLYCDAATNRN	299
OY	298	FARLDVFLFLPPSGGMORTRVYGVGSNPMWNSAVCVYSLGIDRDYRFTSLSKGYHMGS	357
Db	300	FARLDVDFLLPPSGGMORTRVYGVGSNPMWNSAVCVYSLSGLDIDKDYFRITSLSKGYHSLP	359
OY	358	NRPGMCLPRKOPITETFOVADSHPEVAQRVEPMGR	394

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Db      360 NRPFGKCLPDDQPIPTETFOVADRHPEVAQRVEPMGP    396
                                                                    ||||| ||| :||||| ||||| ||||| ||||| |||||
RESULT 4
ID      AAY28552 standard; Peptide: 666 AA.
XX      AAY28552;
XX      SBSEMWL polypeptide #1.
DE      19-OCT-1999 (first entry)
XX      SBSEMWL polypeptide #1.
XX      SBSEMWL; semaphorin; axon outgrowth; multidrug resistance; spinal injury;
KW      neurodegeneration; viral infection; cancer.
OS      Homo sapiens.
PN      WO9338885-A2.
XX      05-AUG-1999.
PD      25-JAN-1999; 99WO-EP00422.
PF      30-JAN-1998; 98EP-0300694.
PR      (SMIK ) SMITHKLIN BEECHAM PLC.
PA      Hayes PD, Michalovich D;
PI      WPI; 1999-479166/40.
DR      N-PsDB: AAZ00102.
XX      Novel SBSEMWL molecules used for treating neurodegeneration, spinal
PT      injury, neuropathies, and neuromuscular, psychiatric, and
PT      inflammatory disorders, developmental malfunctions, cancer, immune
PT      system disorders and viral infections
PS      Claim 4; Page 29-31; 34pp; English.
XX      This sequence is human SBSEMWL polypeptide #1. SBSEMWL polypeptides are
CC      believed to be members of the Semaphorin family of polypeptides.
CC      Semaphorin polypeptides act as recognition molecules and are involved in
CC      axon outgrowth control. They are also likely to have a role in immune
CC      function and multidrug resistance. SBSEMWL polypeptides may be used for
CC      detecting diseases associated with inappropriate SBSEMWL activity or
CC      levels. SBSEMWL polypeptides and polynucleotides, agonists, antagonists
CC      and antibodies are used to treat neurodegeneration, spinal injury,
CC      neuropathies, and neuromuscular, psychiatric, and inflammatory disorders,
CC      developmental malfunctions, cancer, disorders of the immune system and
CC      viral infection. The polynucleotide is also useful as a source of primers
CC      and probes, and also for detecting the above diseases.
SQ      Sequence        666 AA:
Query Match          89.5%; Score 1897; DB 20; Length 666;
Best Local Similarity 90.2%; Pred. No. 3,3e+181;
Matches 358; Conservative 8; Mismatches 27; Indels 4; Gaps 2
1 MTPPPGGAAPSAAPARVALSLPARFGLPLRLILLFWVAASAOGRSHSGPRISAVMK- 59
  | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db      1 MTPPPGGAAPSAAPARVALSPPARLGLPLRLRLLILLMAAASAOGRHLSGPRIFAWMG 60
  | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
QY      60 --GQDHVPDSQDEPHTVLFHEPGSSVWVGGRKYHNENPEGKNA SVTVNIGSTKGC 117
  | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db      61 HVGODRVDPGGQTEPHVLFEHPGSSVWVGGRKYLVLPDFPGKNASVTVNIGSTKGC 120
  | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
QY      118 QKKDQGNITTLERNGNLVCGTAAKRPCSCMNLYNDVSILGEMKGTA PPSDENSL 177
  | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db      121 LDKRCENTITLLERSSEGLACGTAAHPSCMNLVNGTV--PLGMRGTAFPSDENSL 179
  | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
QY      178 VLEEGEVSVSTRKQENKIRFRRIRESELSYSDTWQNQPOFIKATIVHQDAQYDK 237

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Db 180 VLFEGDEVYSTIRKQYENCKIPRFRIRIGESSELYSDTVMNPQFIKATIVHQDAYDOK 239
QY 238 IYFFREDNPDKNPAPLNVSRVAQLCRDQGESSELSVSKNFTLKAMLVCSDAATNRN 297
Db 240 IYFFREDNPDKNPAPLNVSRVAQLCRDQGESSELSVSKNFTLKAMLVCSDAATNRN 299
QY 238 FNRLODVFLLPDPSCQMDRTRYGVFSNPMWNSAVCVYSLGIDIVFRTSSLKGYHMGSL 357
Db 300 FNRLODVFLLPDPSCQMDRTRYGVFSNPMWNSAVCVYSLGIDIVFRTSSLKGYHSSLP 359
QY 358 NRPFGMCLPKKOPITETFOVADSHPEVAQRVEPMGP 394
Db 360 NRPFGKCLPDQOPIPTETFOVADRHPEVAQRVEPMGP 396

RESULT 5
AAW92804
ID AAW92804 standard; Protein: 666 AA.
XX
AC AAW92804;
XX
DT 07-MAY-1999 (first entry)
XX
DE EP-892047 Seq ID 3.
XX
KW Semaphorin L; human; immunosuppressant; anti-inflammatory; gene therapy;
KW organ transplantation; inflammation therapy; immunotherapy; agonist;
KW immunomodulatory; antagonist.
XX
OS Homo sapiens.
XX
PN EP892047-A2.
XX
PD 20-JAN-1999.
XX
PE 06-JUL-1998; 98EP-0112470.
XX
PR 11-FEB-1998; 98DE-1005371.
XX
PR 09-JUL-1997; 97DE-1029211.
XX
PA (HMRI ) HOECHST MARION ROUSSEL DEUT GMBH.
XX
PI Ensser A, Fleckenstein B;
XX
DR WPI: 1999-083564/08.
XX
PT New semaphorin L proteins - used as immunosuppressants and
PT antiinflammatory agents in organ transplants, inflammation therapy,
PT immunotherapy and gene therapy
XX
PS Claim 2; Page 61-64; 135pp; German.
XX
CC This invention describes a novel human semaphorin L protein. This protein
CC or its encoding DNA are useful as immunosuppressants and/or
CC anti-inflammatory agents in organ transplantation, inflammation therapy,
CC immunotherapy and gene therapy. The DNA can be used to produce knock-out
CC or knock-in animals for research purposes. The proteins or DNA can be
CC used to search for the corresponding receptors or to screen for
CC immunomodulatory agonists or antagonists.
XX
SQ Sequence 666 AA;

Query Match 89.5%; Score 1897; DB 20; Length 666;
Best Local Similarity 90.2%; Pred. No. 3.3e-181;
Matches 358; Conservative 8; Mismatches 27; Indels 4; Gaps 2;

QY 1 MTPPPGRRAPASAPRARVLSIPAREGLPLRLRLLLVFWVAASAOGHSRSGPRISAWK- 59
Db 1 MTPPPGRRAPASAPRARVLSIPAREGLPLRLRLLLVFWVAASAOGHSRSGPRIFAQWK 60
QY 60 --GDDHDFQSPREHTVLFHFGSFSVWVGGRGKVYHFNPEEGKNASVRYTNIQSTGSC 117
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Db 61 HVGQDRVDFGQTEPHTVLFHEBGSSSVWVGGRGKYLLFDPEEGKNASVRYTNIQSTGSC 120
QY 118 ODKQDCGNITTLERRGNGLVCGTNARKPSCMNLYNDVSVSLGPMKGAFSPDENSL 177
Db 121 LDKRCRCENTITTLERRSEBLLACGTNARHPSCMNLYNGIV-PLGEMRGYAFSPDENSL 179
QY 178 VLFEGDEVYSTIRKQYENCKIPRFRIRIGESSELYSDTVMNPQFIKATIVHQDAYDOK 237
Db 180 VLFEGDEVYSTIRKQYENCKIPRFRIRIGESSELYSDTVMNPQFIKATIVHQDAYDOK 239
QY 238 IYFFREDNPDKNPAPLNVSRVAQLCRDQGESSELSVSKNFTLKAMLVCSDAATNRN 297
Db 240 IYFFREDNPDKNPAPLNVSRVAQLCRDQGESSELSVSKNFTLKAMLVCSDAATNRN 299
QY 238 FNRLODVFLLPDPSCQMDRTRYGVFSNPMWNSAVCVYSLGIDIVFRTSSLKGYHMGSL 357
Db 300 FNRLODVFLLPDPSCQMDRTRYGVFSNPMWNSAVCVYSLGIDIVFRTSSLKGYHSSLP 359
QY 358 NRPFGMCLPKKOPITETFOVADSHPEVAQRVEPMGP 394
Db 360 NRPFGKCLPDQOPIPTETFOVADRHPEVAQRVEPMGP 396
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RESULT 6
AAV82433
ID AAV82433 standard; Protein: 666 AA.
XX
AC AAV82433;
XX
DT 27-JUN-2000 (first entry)
XX
DE Human CDw108 protein SEQ ID NO:1.
XX
KW Human; CDw108; detection; diagnosis; HIV; infection; anti-HIV.
XX
OS Homo sapiens.
XX
PN WO200012700-A1.
XX
PD 09-MAR-2000.
XX
PE 25-AUG-1999; 99WO-JP04571.
XX
PR 26-AUG-1998; 98JP-0239687.
XX
PA (SHIO ) SHIONOGI & CO LTD.
XX
PI Yamada A, Kubo K, Itoh K;
XX
DR WPI: 2000-246752/21.
XX
DR N-PSDB; AAA08183.
XX
PT New CDw108 protein, useful in diagnosis of and as remedy for
PT CDw108-associated diseases e.g. HIV-1 infection, and in study of
PT biological functions and molecular specificity of CDw108
XX
PS Claim 1; Fig 1; 73pp; Japanese.
XX
CC The present sequence represents human CDw108. The CDw109 nucleotide
CC and protein sequences can be used in the diagnosis and treatment of
CC CDw108-associated diseases e.g. HIV-1 infection, and in study of
CC biological functions and molecular specificity of CDw108.
XX
SQ Sequence 666 AA;

Query Match 89.5%; Score 1897; DB 21; Length 666;
Best Local Similarity 90.2%; Pred. No. 3.3e-181;
Matches 358; Conservative 8; Mismatches 27; Indels 4; Gaps 2;

QY 1 MTPPPGRRAPASAPRARVLSIPAREGLPLRLRLLLVFWVAASAOGHSRSGPRISAWK- 59
Db 1 MTPPPGRRAPASAPRARVLSIPAREGLPLRLRLLLVFWVAASAOGHSRSGPRIFAQWK 60
QY 60 --GDDHDFQSPREHTVLFHFGSFSVWVGGRGKVYHFNPEEGKNASVRYTNIQSTGSC 117
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XX Use of Sema7A polypeptides, polynucleotides, or activators or  
 PT inhibitors, for manufacturing a medicament for treating autoimmune  
 PT disorders, chronic obstructive pulmonary disease, inflammation,  
 PT psoriasis and stroke -  
 PS Claim 5; Page 28-29; 36pp; English.  
 XX  
 CC The present sequence is human Sema7A protein which  
 CC belongs to semaphorin family. The Sema7A gene is located on human  
 CC chromosome 15q22.3-q23. The Sema7A polypeptide and polynucleotide  
 CC are useful for the manufacture of a medicament for treating autoimmune  
 CC disorders, chronic obstructive pulmonary disease (COPD), inflammation,  
 CC psoriasis, tissue repair, wounds, to enhance wound healing, irritable  
 CC bowel syndrome, stroke, atherosclerosis, cancer or diseases where it is  
 CC necessary to promote dendritic cell formation. The polypeptide is also  
 CC useful in the diagnostic assays for detecting diseases associated with  
 CC inappropriate levels or activities of Sema7A.  
 XX  
 SQ Sequence 666 AA:  
 Query Match 89.5%; Score 1897; DB 22; Length 666;  
 Best Local Similarity 90.2%; Pred. No. 3.3e-181;  
 Matches 358; Conservative 8; Mismatches 27; Indels 4; Gaps 2;  
 QY 1 MTPPPGGAAPSAPRARVLSLPARGCLPLRLLLVFWAASAOCHSRSGPRISAVWK- 59  
 DB 1 MTPPPGGAAPSAPRARVGPAPRLGLPLRLLLMAAASAOCHLRSGPRIFAVWKG 60  
 QY 60 --GODHVDSEPHPTVLFHEPGSFVWVGKGYVHFNPPGKNASVTVMNGSTKGC 117  
 DB 61 HVGODRVDFGQTEPHTVLFHEPGSSVWVGKGYVLFDPGKKNASVTVMNGSTKGC 120  
 QY 118 ODKODCGNYITLLERKGNCLVCGTNARKPSQMNLVNDSVMSLGEKGYAFSPDENS 177  
 DB 121 LDKRDCENTYTLLEKRSSEGLACGTNARKPSQMNLVNDSVMSLGEKGYAFSPDENS 179  
 QY 178 VLFEGDEVYSTIRKQETNKKIRPRIRIGESSELYTSQYVWQNPQFKATIVHODAYDDK 237  
 DB 180 VLFEGDEVYSTIRKQETNKKIRPRIRIGESSELYTSQYVWQNPQFKATIVHODAYDDK 239  
 QY 238 IYFFREDPDKNPEAPLVNRSVAQLCRGDGESSLSVSKNTFLKALVCSDAATNEN 297  
 DB 240 IYFFREDPDKNPEAPLVNRSVAQLCRGDGESSLSVSKNTFLKALVCSDAATNEN 299  
 QY 298 FNRLODVELLPDPSQWMDTRVYGVFSNPMNYSACVYSLGIDIRVFTSSLKGYHMGSL 357  
 DB 300 FNRLODVELLPDPSQWMDTRVYGVFSNPMNYSACVYSLGIDIRVFTSSLKGYHSSLR 359  
 QY 358 NRPFGMCLPKKQPIPTETFGVADSHPEVAQRYEPMGP 394  
 DB 360 NRPFGMCLPKKQPIPTETFGVADSHPEVAQRYEPMGP 396  
 RESULT 9  
 AAY56854  
 ID AAY56854 standard; Protein; 634 AA.  
 AC AAY56854;  
 XX  
 DT 10-APR-2000 (first entry)  
 XX  
 DE Human semaphorin K1 polypeptide.  
 XX  
 KW Semaphorin K1; cellular physiology; neurite outgrowth; neuron; human;  
 KW immunogen; pharmaceutical.  
 XX  
 OS Homo sapiens.  
 XX  
 XX JP1341988-A.  
 XX  
 PD 14-DEC-1999.  
 XX

PF 11-MAR-1999; 99JP-0065672.  
 XX  
 PR 11-MAR-1998; 98US-0041236.  
 XX  
 PA (EXEL-) EXELIXIS PHARM INC.  
 XX  
 DR WPI: 2000-109378/10.  
 DR N-PSDB: AAZ46841.  
 XX  
 PT New semaphorin polypeptides, useful cell physiology modulators and  
 PT immunogens -  
 PS Claim 1; Page 12-15; 57pp; Japanese.  
 XX  
 CC The invention provided isolated human semaphorin K1 polypeptides. The  
 CC polypeptides, or nucleic acids encoding them, can be used to modulate  
 CC cellular physiology by modulating semaphorin K1 activity, e.g. semaphorin  
 CC K1 polypeptide fragments or antisense nucleic acids can be used to  
 CC enhance neurite outgrowth from damaged neurons. The polypeptides can also  
 CC be used as immunogens, reagents for isolating other semaphorins, or as  
 CC reagents for screening chemical libraries for lead pharmaceutical agents.  
 CC The nucleic acids can also be used as probes and primers for diagnostic  
 CC purposes. The present sequence represents the human semaphorin K1  
 CC polypeptide.  
 XX  
 SQ Sequence 634 AA:  
 Query Match 82.8%; Score 1755; DB 21; Length 634;  
 Best Local Similarity 90.4%; Pred. No. 5.7e-167;  
 Matches 330; Conservative 8; Mismatches 23; Indels 4; Gaps 2;  
 QY 33 LLLVFWAASAOCHSRSGPRISAVWK---GODHVDSEPHPTVLFHEPGSFVWVGR 89  
 DB 1 LLLVFWAASAOCHLRSGPRIFAVWKGHVGDQVDFGQTEPHTVLFHEPGSSVWVGR 60  
 QY 90 GKVYHFNPEPGKNASVTVMNGSTKGCODKDCGNYITLLERKGNCLVCGTNARKPSC 149  
 DB 61 GKVYHFNPEPGKNASVTVMNGSTKGCODKDCGNYITLLERKGNCLVCGTNARKPSC 120  
 QY 150 WNLVNDVMSLGEKGYAFSPDENSILVLFEGDEVYSTIRKQETNKKIRPRIRIGESE 209  
 DB 121 WNLVNTVY-PLGEMKGYAFSPDENSILVLFEGDEVYSTIRKQETNKKIRPRIRIGESE 179  
 QY 210 LYTSDTVMQNPQFKATIVHODAYDDKIYFFREDPDKNPEAPLVNRSVAQLCRGDG 269  
 DB 180 LYTSDTVMQNPQFKATIVHODAYDDKIYFFREDPDKNPEAPLVNRSVAQLCRGDG 239  
 QY 270 GESSLSVSKNMFELKAMLVCSDAATNENRLODVELLPDPSQWMDTRVYGVFSNPMNY 329  
 DB 240 GESSLSVSKNMFELKAMLVCSDAATNENRLODVELLPDPSQWMDTRVYGVFSNPMNY 299  
 QY 330 SAVCYSLGIDIRVFTSSLKGYHMGSLNRPFGMCLPKKQPIPTETFGVADSHPEVAQRY 389  
 DB 300 SAVCYSLGIDIRVFTSSLKGYHSSLRNPFGMCLPKKQPIPTETFGVADSHPEVAQRY 359  
 QY 390 EPMGP 394  
 DB 360 EPMGP 364  
 RESULT 10  
 AAY28553  
 ID AAY28553 standard; Peptide; 215 AA.  
 AC AAY28553;  
 XX  
 DT 19-OCT-1999 (first entry)  
 XX  
 DE SBSEMWL polypeptide #2.  
 XX  
 KW SBSEMWL; semaphorin; axon outgrowth; multidrug resistance; spinal injury;  
 KW neurodegeneration; viral infection; cancer.  
 XX

OS Homo sapiens.  
 XX  
 PN W09938885-A2.  
 XX  
 PD 05-AUG-1999.  
 XX  
 PF 25-JAN-1999; 99WO-EP00422.  
 XX  
 PR 30-JAN-1998; 98EP-0300694.  
 XX  
 PA (SMIK ) SMITHKLINE BEECHAM PLC.  
 PI  
 PI Hayes PD, Michalovich D;  
 XX  
 DR WPI: 1999-479166/40.  
 DR N-PSDB; AAZ00103.  
 XX  
 PT Novel SBSEMWL molecules used for treating neurodegeneration, spinal  
 PT injury, neuropathies, and neuromuscular, psychiatric, and  
 PT inflammatory disorders, developmental malfunctions, cancer, immune  
 PT system disorders and viral infections  
 XX  
 PS Claim 17; Page 32; 34pp; English.  
 XX  
 CC This sequence is human SBSEMWL polypeptide #2. SBSEMWL polypeptides are  
 CC believed to be members of the Semaphorin family of polypeptides.  
 CC Semaphorin polypeptides act as recognition molecules and are involved in  
 CC axon outgrowth control. They are also likely to have a role in immune  
 CC function and multidrug resistance. SBSEMWL polypeptides may be used for  
 CC detecting diseases associated with inappropriate SBSEMWL activity or  
 CC levels. SBSEMWL polypeptides and polynucleotides, agonists, antagonists  
 CC and antibodies are used to treat neurodegeneration, spinal injury,  
 CC neuropathies, and neuromuscular, psychiatric, and inflammatory disorders,  
 CC developmental malfunctions, cancer, disorders of the immune system and  
 CC viral infection. The polynucleotide is also useful as a source of primers  
 CC and probes, and also for detecting the above diseases.  
 XX  
 SQ Sequence 215 AA;  
 XX  
 Query Match 44.7%; Score 948.5; DB 20; Length 215;  
 Best Local Similarity 85.2%; Pred. No. 11e-86;  
 Matches 184; Conservative 6; Mismatches 23; Indels 3; Gaps 3;  
 XX  
 OY 60 GODHDFSEPHPTVLFHPGGSFVWVGGRGVYHFPPEGKNASVRYTNIGSTKSCOD 119  
 DB 1 GQDRVDFGTEPHVYLFHPGSSVWVGGRGVYLFDFEGKNASVRYTNIGSTKSCOD 60  
 OY 120 KODCGNYITLERRNGLLVCGTNAKPPSCWNLVNDVSVSLGEMKGYAPFSPDEN-SLV 178  
 DB 61 KRDCENYITLERRSEGLACGTNAHPPSCWNLVN-ALMCHLGESGAYAPFSPDENVPWF 119  
 OY 179 LFEQGEVYSTIRK-QEYNKIRFRIRIRGESELYTSDTYWQNPQFIKATIVHODQAYDK 237  
 DB 120 CEQGEVYSTIRKARVYNNEDPRFRIRIRGESELYTSDTYWQNPQFIKATIVHODQAYDK 179  
 OY 238 IYFFREDNPDKNPEAPLVNSRYAOLCRGDGESS 273  
 DB 180 IYFFREDNPDKNPEAPLVNSRYAOLCRGDGESS 215  
 XX  
 RESULT 11  
 AAY56855  
 ID AAY56855 standard; Protein: 606 AA.  
 XX  
 AC AAY56855;  
 XX  
 DT 10-APR-2000 (first entry)  
 XX  
 DE Semaphorin K1 polypeptide related sequence.  
 XX  
 KW Semaphorin K1, cellular physiology; neurite outgrowth; neuron; human;  
 XX immunogen; pharmaceutical.  
 XX

OS Unidentified.  
 XX  
 PN JP11341988-A.  
 XX  
 PD 14-DEC-1999.  
 XX  
 PF 11-MAR-1999; 99JP-0065672.  
 XX  
 PR 11-MAR-1998; 98US-0041236.  
 XX  
 PA (EXEL-) EXELIXIS PHARM INC.  
 PI  
 PI WPI: 2000-109378/10.  
 DR N-PSDB; AAZ46842.  
 XX  
 PT New semaphorin polypeptides, useful cell physiology modulators and  
 PT immunogens -  
 XX  
 PS Disclosure; Page 17-20; 57pp; Japanese.  
 XX  
 CC The invention provided isolated human semaphorin K1 polypeptides. The  
 CC polypeptides, or nucleic acids encoding them, can be used to modulate  
 CC cellular physiology by modulating semaphorin K1 activity, e.g. semaphorin  
 CC K1 polypeptide fragments or antisense nucleic acids can be used to  
 CC enhance neurite outgrowth from damaged neurons. The polypeptides can also  
 CC be used as immunogens, reagents for isolating other semaphorins, or as  
 CC reagents for screening chemical libraries for lead pharmaceutical agents.  
 CC The nucleic acids can also be used as probes and primers for diagnostic  
 CC purposes.  
 XX  
 SQ Sequence 606 AA;  
 XX  
 Query Match 42.5%; Score 902; DB 21; Length 606;  
 Best Local Similarity 48.9%; Pred. No. 2.7e-81;  
 Matches 179; Conservative 54; Mismatches 127; Indels 6; Gaps 4;  
 XX  
 OY 27 LPLRLRLLVFWAASAOGHSRSGPRISAVMK---GODHDFSEPHPTVLFHPGGSF 83  
 DB 4 LCVSIRLLMIL-SAITAASRFIDKPRLLVNLTDGRGQ-HRFGQPEPHVLFHSLNSTD 61  
 OY 84 VVVGGRGVYHFPPEGKNASVRYTNIGSTKSCODKODCGNYITLERRNGLLVCGTN 143  
 DB 62 VVVGGNNTIYLFDFAHSSNASTALINITSTHNRHLSGCCENFTLLHNOTDGLACGTN 121  
 OY 144 ARKPSQWNLVNDVSVSLGEMKGYAPFSPDENLVLEQGEVYSTIRKQYNNKIRFR 203  
 DB 122 SQRPSQW-LINNLTQFLGPKGLAPFSPSSGNLVLFDDNDYSTINLYKSLSGSHKFR 180  
 OY 204 IRGESELYTSDTYWQNPQFIKATIVHODQAYDKIYFFREDNPDKNPEAPLVNSRYAOL 263  
 DB 181 IAGQVELYTSIDTAMHRPQVQATAVHKNESYDKIITFFQENSHSDFKOPRHPVPRVGV 240  
 OY 264 CRGDGESSLSVSKNNTFLKAMLVCSDAATNENRDLDFLLPPSGQWRDTRYGVF 323  
 DB 241 CSSDQGESLSVYKNTFLKARLACVDYDGRIVNELDIFIQAPENSMEETLLIYGLF 300  
 OY 324 SNPNWNSAVCVYSLGDIDRVFRTSSLSKGYHMGSLNRPQCLRKQPIPETEQVADSHR 383  
 DB 301 LSPWNFSAVCVFTVKDIDHVEFKTSKLKNYHKLPTPRPGCCMKNHGVPTETEQVADRYP 360  
 OY 384 EVAQRV 389  
 DB 361 EVADPV 366  
 XX  
 RESULT 12  
 AAB93440  
 ID AAB93440 standard; Protein: 379 AA.  
 XX  
 AC AAB93440;  
 XX  
 DT 26-JUN-2001 (first entry)  
 XX  
 DE  
 XX  
 KW  
 XX immunogen;  
 XX

DE Human protein sequence SEQ ID NO:12678.  
XX  
KW Human; primer; detection; diagnosis; antisense therapy; gene therapy.  
XX  
OS Homo sapiens.  
XX EPI074617-A2.  
XX  
PD 07-FEB-2001.  
XX  
PF 28-JUL-2000; 2000EP-0116126.  
XX  
PR 29-JUL-1999; 99JP-0248036.  
PR 27-AUG-1999; 99JP-0300253.  
PR 11-JAN-2000; 2000JP-0118776.  
PR 02-MAY-2000; 2000JP-0183767.  
PR 09-JUN-2000; 2000JP-0241899.  
XX  
PA (HELI-) HELIX RES INST.  
XX  
PI Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;  
PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;  
XX  
DR WPI: 2001-318749/34.  
XX  
PT Primer sets for synthesizing polynucleotides, particularly the 5602  
PT full-length cDNAs defined in the specification, and for the detection  
PT and/or diagnosis of the abnormality of the proteins encoded by the  
PT full-length cDNAs -  
XX  
PS Claim 8; SEQ ID 12678; 2537bp + CD ROM; English.  
XX  
CC The present invention describes primer sets for synthesizing 5602  
CC full-length cDNAs defined in the specification. Where a primer set  
CC comprises: (a) an oligo-dT primer and an oligonucleotide complementary  
CC to the complementary strand of a polynucleotide which comprises one of  
CC the 5602 nucleotide sequences defined in the specification, where the  
CC oligonucleotide comprises at least 15 nucleotides; or (b) a combination  
CC of an oligonucleotide comprising a sequence complementary to the  
CC complementary strand of a polynucleotide which comprises a 5'-end  
CC sequence and an oligonucleotide comprising a sequence complementary to a  
CC polynucleotide which comprises a 3'-end sequence, where the  
CC oligonucleotide comprises at least 15 nucleotides and the combination of  
CC the 5'-end sequence/3'-end sequence is selected from those defined in  
CC the specification. The primer sets can be used in antisense therapy and  
CC in gene therapy. The primers are useful for synthesizing polynucleotides,  
CC particularly full-length cDNAs. The primers are also useful for the  
CC detection and/or diagnosis of the abnormality of the proteins encoded by  
CC the full-length cDNAs. The primers allow obtaining of the full-length  
CC cDNAs easily without any specialised methods. AAH03166 to AAH13628 and  
CC AAH13633 to AAH18742 represent human cDNA sequences; AAB92446 to  
CC AAB95893 represent human amino acid sequences; and AAH13629 to AAH13632  
CC represent oligonucleotides, all of which are used in the exemplification  
CC of the present invention.  
XX  
SQ Sequence 379 AA:  
Query Match 26.0%; Score 552; DB 22; Length 379;  
Best Local Similarity 91.7%; Pred. No. 1.9e-46;  
Matches 100; Conservative 3; Mismatches 6; Indels 0; Gaps 0;  
QY 286 MLVCSDATNNFNRLQOVFLPPSGGWRDTRRYGVFSNPNWNSAVCVYSGLDIDRFR 345  
DB 1 MLVCSDATNNFNRLQOVFLPPSGGWRDTRRYGVFSNPNWNSAVCVYSGLDIDRFR 60  
QY 346 TTSKGYMGSLNPPGMCPLPKKQPIPTETFOVADSHPEVAVQRYVPMGP 394  
DB 61 TTSKGYMGSLNPPGMCPLPKKQPIPTETFOVADSHPEVAVQRYVPMGP 109

XX ABB97964;  
AC 06-SEP-2002 (first entry)  
XX  
DT Human protein sequence #31.  
XX  
DE Human; brain; tonsil; hippocampus; foetal brain; diagnosis.  
XX  
KW Homo sapiens.  
XX  
OS WO200252005-A1.  
XX  
PN 04-JUL-2002.  
XX  
PD 20-DEC-2001; 2001WO-JP11217.  
XX  
PF 22-DEC-2000; 2000JP-0389742.  
XX  
PR (KAZU-) KAZUSA DNA RES INST FOUND.  
XX (CELE-) CELESTAR LEXICO-SCI LTD.  
XX  
PI Ohara O, Nagase T, Nakajima D;  
XX  
DR WPI: 2002-500762/53.  
DR N-PSDB: ABN83984.  
XX  
PT Genes and their expression products cloned from human cDNA libraries  
PT for treatment and diagnosis of diseases associated with their  
PT expression -  
XX  
PS Claim 1(a); Page 126-132; 238bp; Japanese.  
XX  
CC The invention relates to DNA encoding polypeptides directly cloned from  
CC cDNA libraries originating in adult whole brain, human tonsil, human  
CC adult hippocampus and human foetal whole brain. Polypeptides and  
CC polynucleotides of the invention may be used in the investigation of  
CC differential expression of the DNA sequences in normal subjects and  
CC disease patients. They may also be used in the production of antibodies,  
CC oligonucleotide probes and DNA chips for diagnosis and identification  
CC of drugs for treatment of diseases with which the DNA sequences are  
CC associated. The sequences given in records ABB97934-ABB97964 represent  
CC human proteins of the invention.  
XX  
SQ Sequence 893 AA:  
Query Match 18.7%; Score 397; DB 23; Length 893;  
Best Local Similarity 30.1%; Pred. No. 2.7e-30;  
Matches 129; Conservative 54; Mismatches 167; Indels 78; Gaps 16;  
QY 3 PPPGCAAPASAPRAVFL-----SLPARFGPLRLRLLVF-----WVAA 41  
DB 42 PPEPEPDTVA PALMLRTAMGLRSM LAAPWGA LPPRPPLLLLLLLLP PPTWALS 101  
QY 42 ASAGSHSRGPRISAVMKGDHVDSPQEP-----HTVLFHFGSFSWVGRCGVYHFN 96  
DB 102 -----PRIS-LPUGSEPPFLREFAEHISVYTA LLSRDGRTYVAGREALFALS 150  
QY 97 -----FPEGKNAVSVTVIGSTKGSCKODK-----ODCNYI-TLLERGNGLVCGTNAR 145  
DB 151 SNLSLPLPGGEVQELLMGADAEKKQCSFKGDPQDCQNYIKILPLSGSHLFTCGTAAAF 210  
QY 146 KPSCW--NLVNDVV-----MSIGEMKVAPFSFDENSLVLFEGDEVYSTIRKQETNGK 197  
DB 211 SPMCTYIMNENFTLARDEKGNVLEDDGRCRPFDPNFKSTALVYNGELY-TGVASSFGGN 269  
QY 198 IPRFRIRIGSEELITSQV--MNPQFIKATIVHD-----QAYDDKITYFPREDNPKMP 251  
DB 270 DPAISRQSILRPPTKTESSLNMLQDPAFVASAYIPISLGSGLGDDDKIYFFFSSETQEEEF 329  
QY 252 EAPLVNVRVAOLCRDQGESLSVSKNWFPLKALVCSDAATNNFNRLQOVFLPPPS 311  
DB 330 FENTIVSRILARICKDGEGERVLQ-QRWTSFLKQOLLCSRPDDGPFVNLQDVFTLSFSP 388



OY	312	GOMDTRRYGFGFNSPMWN-----SAYCVYISLGDIDRF-----RTSLKGYHMGSL	357
Db	389	QDMRDTELYGFTSQWMHGTTGSAVCFYTKMDQVRVFSGLYKVENRTEQMYTVTHHPV	448
OY	358	NPRGMCL 365	
Db	449	TPRGACI 456	
RESULT 14			
ID	AAW17658		
AC	AAW17658	standard; Protein: 861 AA.	
XX	AAW17658;		
XX	24-JUL-1997	(first entry)	
DE	Mouse CD100 antigen.		
KW	CD100 antigen; semaphorin; leukocyte; B cell; T cell; lymphocyte; vaccine.		
XX	Mus sp.		
XX	Key	Location/Qualifiers	
FT	Peptide	1..41	
FT		/label= sig_peptide	
FT	Protein	42..861	
FT		/label= Mat_protein	
FT	Domain	42..553	
FT		/label= Semaphorin_domain	
FT	Domain	554..630	
FT		/label= Ig-like_domain	
FT	Domain	631..732	
FT		/label= stalk_domain	
FT	Domain	734..752	
FT		/label= Transmembrane_domain	
FT	Domain	753..861	
FT		/label= Cytoplasmic_domain	
FT	Modified-site	807..814	
FT		/label= Phosphorylation	
XX		/note= "putative tyrosine phosphorylation site"	
PN	W09717368-A1.		
PD	15-MAY-1997.		
PF	12-NOV-1996;	96WO-US18645.	
XX	09-NOV-1995;	95US-0556422.	
PA	(DAND ) DANA FARBER CANCER INST.		
PI	Boussiotis V, Freeman GJ, Hall KT, Nadler LM, Schultze JL;		
DR	WPI: 1997-280982/25.		
XX	N-PSDB; AAT60666.		
XX	Nucleic acid molecule encoding CD100 antigen - which stimulates leukocyte response, e.g. B cell aggregation, differentiation, survival and T cell proliferation		
PS	Example 8; Page 86-89; 135pp; English.		
XX	Mouse CD100 antigen (AAW17657) is a novel leukocyte semaphorin-like protein that stimulates a leukocyte response, including B cell aggregation, B cell differentiation, B cell survival and/or T cell proliferation. Its amino acid sequence was deduced from a cDNA clone (AAT60666) isolated from murine T cells. Human CD100 antigen (AAW17657) has also been identified. CD100 polypeptides and fusion proteins, nucleic acids, and host cells expressing CD100 can be utilized in diagnostic and therapeutic methods involving modulation		

[illegible]

